Amendments To The Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

In the Claims:

What is claimed is:

1. (Currently amended) A compound of formula (I):

$$R_1$$
 Z
 $(R^4)_r$
 $(CH_2)_m$
 O
 R^3

wherein:

R¹ represents phenyl which may be optionally substituted by one or more substituents which may be the same or different and which are selected from the group consisting of: halogen; trifluoromethyl; -C₁₋₆alkyl optionally substituted by COOR¹⁵; -C₁₋₆alkoxy optionally substituted by COOR¹⁵; hydroxy; oxo; cyano; -C₁₋₆alkyl-cyano; C₂₋₆alkenyl optionally substituted by COOR¹⁵; C₃₋₇cycloalkyl; C₁₋₆alkylsulfonyl; C₂₋₆alkenoxy C₁₋₆alkylthio; NR¹⁵R¹⁶; -C₁₋₆alkyl-aryl; aryl; -CO-aryl optionally substituted by halogen; -CO-heteroaryl; -CO-heterocyclyl; -COOR¹⁵; -COR¹⁵; -CONR¹⁵R¹⁶ optionally substituted by C₁₋₆alkyl, halogen or -C₁₋₆alkylC₁₋₆alkoxy; and -C₁₋₆alkyl-CO-aryl groups; and in which

 R^{15} and R^{16} independently represent hydrogen, C_{1-6} alkyl or C_{3-8} cycloalkyl or together may be fused to form a 5- to 7-membered non-aromatic heterocyclic ring optionally interrupted by an O or S atom and optionally substituted by a halogen, C_{1-6} alkyl or C_{1-6} alkyl C_{1-6} alkoxy group;

Z represents CO;

r is 0;

p is 1;

m is 0;

R³ represents <u>a group of formula (i):</u>

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$$-(CH_2)_f$$
 $(R^{14})_k$ (i)

wherein

f is 0;

g is 2;

h is 1;

k is 0; and

 R^{13} represents C_{1-6} alkyl or C_{3-8} cycloalkyl;

or a pharmaceutically acceptable salt thereof.

2-11. (Cancelled)

Add the following new claims:

12. (Currently amended)

A compound according to claim 1 wherein R¹ is phenyl which may be optionally substituted by 1, 2 or 3 substituents which may be the same or different and which are selected from the group consisting of: chlorine, fluorine, bromine; trifluoromethyl; methyl, ethyl, isopropyl, propyl, t-butyl (optionally substituted by COOH, COOMe or COOEt); methoxy, butoxy, -OCH(Me)₂, -OC(Me)₃ (optionally substituted by COOH or COOMe); hydroxy; oxo; cyano; -CH₂-CN; ethenyl (optionally substituted by COOMe); cyclopentyl; -SO₂Me; -OCH₂CH=CH₂; -S-ethyl; N(Me)₂; benzyl; phenyl; -CO-phenyl (optionally substituted by chlorine); -CO-azetidinyl; -CO-tetrahydropyranyl; COOH, COOMe, COOt-butyl; -CO-methyl, -CO-ethyl, -CO-isopropyl, -CO-cyclopropyl, -CO-cyclobutyl, -CO-cyclopentyl, -CO-cyclohexyl; -CONH₂, -CO-pyrrolidinyl, -CO-morpholinyl, -CO-piperazinyl, -CO-piperidinyl, -CO-thiomorpholinyl (optionally substituted by methyl, fluorine and -CH₂OMe); or -CH₂COphenyl groups;

or a pharmaceutically acceptable salt thereof.

13. (Previously presented) A compound according to claim 1 wherein R^1 is phenyl substituted by C_{1-6} alkylsulfonyl.

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- 14. (Previously presented) A compound according to claim 1 wherein R^1 is phenyl substituted by SO_2Me .
- 15. (Previously presented) A compound according to claim 1 wherein R^1 is phenyl substituted by SO_2Me at the para position.
- 16. (Previously presented) A compound according to claim 1 wherein -O-R³ is present at the para position of the phenyl group with respect to the rest of the compound.
- 17. (Previously presented) A compound according to claim 1 wherein R¹³ represents isopropyl, cyclopropyl or cyclobutyl.
- 18. (Previously presented) A compound according to claim 13, wherein R¹³ represents isopropyl, cyclopropyl or cyclobutyl.
- 19. (Previously presented) A compound according to claim 14, wherein R¹³ represents isopropyl, cyclopropyl or cyclobutyl.
- 20. (Previously presented) A compound which is 1-(4-{[1-(1-methylethyl)-4-piperidinyl]oxy}phenyl)-4-{[4-(methylsulfonyl)phenyl]carbonyl}piperazine or a pharmaceutically acceptable salt thereof.
- 21. (Previously presented) A pharmaceutical composition which comprises a compound of formula (I) as defined in claim 1 or a pharmaceutically acceptable salt thereof, and a pharmaceutically acceptable carrier or excipient.
- 22. (Currently amended) A method of treatment of diseases of the upper respiratory tract which comprises administering to a human host in need thereof an effective amount of a compound of formula (I) as defined in claims 1 or a pharmaceutically acceptable salt thereof.

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23. (Previously presented) A method of treatment according to claim 21 in

which the disease is allergic rhinitis.

24. (Previously presented) A pharmaceutical composition which comprises a

compound of formula (I) as defined in claim 18 or a pharmaceutically

acceptable salt thereof, and a pharmaceutically acceptable carrier or excipient.

25. (Currently amended) A method of treatment of diseases of the upper

respiratory tract which comprises administering to a human host in need

thereof an effective amount of a compound of formula (I) as defined in claims

18 or a pharmaceutically acceptable salt thereof.

26. (Previously presented) A method of treatment according to claim 25 in

which the disease is allergic rhinitis.

27. (Previously presented) A pharmaceutical composition which comprises a

compound of formula (I) as defined in claim 19 or a pharmaceutically

acceptable salt thereof, and a pharmaceutically acceptable carrier or excipient.

28. (Currently amended) A method of treatment of diseases of the upper

respiratory tract which comprises administering to a human host in need

thereof an effective amount of a compound of formula (I) as defined in claims

19 or a pharmaceutically acceptable salt thereof.

29. (Previously presented) A method of treatment according to claim 28 in

which the disease is allergic rhinitis.

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